## WHAT IS CLAIMED IS:

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	THE TO CE MINE AG.
1 Qub	1. A cell transduction vector comprising a vector nucleic acid
20	encoding:
3	a retroviral packaging site;
4	a first viral inhibitor subsequence;
5	a splice donor site subsequence;
6	a splice acceptor site subsequence;
7	a retroviral Rev binding subsequence; and,
8	a promoter subsequence;
8 <b>9</b>	wherein:
Ц	

the first viral inhibitor subsequence is located between the splice donor site subsequence and the splice acceptor site subsequence;

the splice donor site subsequence and the splice acceptor site subsequence permit splicing of the first viral inhibitor subsequence from the vector nucleic acid in the nucleus of a cell; and,

the promoter subsequence is operably linked to the first viral inhibitor subsequence.

- 2. The cell transduction vector of claim 1, wherein the vector nucleic acid further encodes a retroviral Rev binding subsequence, wherein the vector nucleic acid is translocated to the cytoplasm in the presence of a Rev protein, and wherein splicing of the first viral inhibitor sequence is inhibited by Rev.
- 3. The cell transduction vector of claim 2, wherein the retroviral Rev binding subsequence is an HIV/RRE sequence.
- 4. The cell transduction vector of claim 1, wherein the first viral inhibitor comprises a nucleic acid subsequence encoding a ribonuclease selected from the pancreatic RNAse A superfamily.
- 5. The cell transduction vector of claim 1, wherein the first viral inhibitor comprises a nucleic acid subsequence encoding a ribonuclease selected from

the group of ribonucleases consisting of Onconase, modified Onconase, and EDN.

- 6. The cell transduction vector of claim 1, wherein the first viral inhibitor subsequence encodes a transdominant protein selected from the group of transdominant proteins consisting of transdominant Gag, transdominant Tat, and transdominant Rev.
  - 7. The cell transduction vector of claim 1, wherein the vector further comprises a cell binding ligand selected from the group consisting of transferrin, *c-kit* ligand, an interleukin and a cytokine.
  - 8. The cell transduction vector of claim 1, wherein the promoter is selected from the group of promoters consisting of a retroviral LTR promoter, a constitutive promoter, an inducible promoter, a tissue specific promoter, a CMV promoter, a probasin promoter and a tetracycline-responsive promoter.
  - 9. The cell transduction vector of claim 1, wherein the vector further comprises an encephalomyocarditis virus internal ribosome entry site (IRES).
  - 10. The cell transduction vector of claim 1, wherein the vector nucleic acid further encodes a second viral inhibitor.
  - 11. The cell transduction vector of claim 9, wherein the vector nucleic acid further encodes a second viral inhibitor, wherein transcription of the second nucleic acid is controlled by the IRES.
  - 12. The cell transduction vector of claim 1, wherein vector nucleic acid further encodes a multicistronic mRNA with a first open reading frame and a second open reading frame, which multicistronic mRNA comprises an IRES sequence which directs translation of the second open reading frame in a cell.

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1		13.	The cell transduction vector of claim 11, wherein the first open			
2	reading frame encodes a viral inhibitor.					
رطس 1 رطس 1	79	14.	The cell transduction vector of claim 1, wherein the vector			
2	comprises a	retrovir	al particle.			
1		15.	The cell transduction vector of claim 1, wherein the vector			
2	nucleic acid	is packa	aged into an HIV particle in a cell infected by a wild-type HIV.			
15		16.	The cell transduction vector of claim 1, wherein the vector			
2 U	nucleic acid is packaged in a liposome.					
19230195 X	1003	17.	The cell transduction vector of claim 14, wherein the retroviral			
2 0	particle is pseudotyped for transduction into hematopoietic stem cells.					
	comprises a	18. pharma	The cell transduction vector of claim 1, wherein the vector further ceutical excipient.			
1	-	19.	The cell transduction vector of claim 1, wherein the vector			
2	nucleic acid	further	encodes a reporter gene.			
الس 1	DDU)	20.	The cell transduction vector of claim 1, wherein the cell			
2	transduction	vector	is selected from the group of cell transduction vectors consisting of			
3	pBAR, pBA	R-ONC	, pBAR-EDN and conservative modifications thereof.			
1		21.	The cell transduction vector of claim 1, wherein the viral inhibitor			
2	is an oncoge	ne inhi	bitor.			
3		22.	The cell transduction vector of claim 1, wherein the vector further			
4	comprises as	n oncog	ene inhibitor.			

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23.

The cell transduction vector of claim 22, wherein the oncogene

2		inhibitor is a nucleic acid encoding an inhibitor selected from the group of inhibitors				
3		consisting of a antibody which specifically binds a Ras protein and an RNAse.				
				•		
1			24.	The cell transduction vector of claim 22, wherein the oncogene		
2		inhibitor is an	ı RNAs	e from the RNAse A superfamily.		
3			25.	A cell transduction vector comprising a nucleic acid subsequence		
4		encoding an	EDN pr	otein, which subsequence is operably linked to a promoter,		
5		wherein said cell transduction vector inhibits the replication of a retrovirus in a cell				
6 🚍		transduced by	the ce	Il transduction vector.		
6 TO TO TO						
1 🖺	Ç	ゆのう	26.	The cell transduction vector of claim 25, wherein the vector is		
2 💆	Ü	pBAR-EDN,	or a co	enservative modification thereof.		
II						
1 =			27.	The cell transduction vector of claim 25, wherein the cell is a		
2 <u> </u> ↓		CD4+ cell.				
1 N 2 N 2 N 2 N 3 N 3 N 3 N 3 N 3 N 3 N 3						
1 <b>9</b>			28.	The cell transduction vector of claim 25, wherein the cell is a		
2		stem cell.				
1			29.	The cell transduction vector of claim 25, wherein the vector		
2		inhibits the re	eplicatio	on of HIV in the cell.		
			-			
1			30.	The cell transduction vector of claim 25, wherein the vector		
2		nucleic acid	is packa	iged in a retroviral particle.		
			.•	•		
1			31.	The cell transduction vector of claim 25, wherein the vector is		
2		packaged in	a liposo			
_		A ··· · · · · · · · · · · · · · · · · ·	¥ - 3 -			
1			32.	The cell transduction vector of claim 25, wherein the vector		
2		comprises a	cell bine	ding ligand selected from the group of cell binding ligands		

consisting of transferrin, kit-ligand, an interleukin, and a cytokine.

			•			
1		33.	The cell transduction vector of claim 25, wherein the vector			
2	nucleic acid f	further (	encodes a subsequence encoding a retroviral chromosome			
3	integration subsequence.					
4 0.1	EL	34.	The dell transduction vector of claim 25, wherein the vector			
5 XW			multicistronic mRNA which encodes a first open reading frame and			
6	a second open reading frame, which multicistronic mRNA is operably linked to a					
7	promoter, wherein the dicistronic mRNA comprises a subsequence encoding EDN.					
	•					
1 🚍		35.	The cell transduction vector of claim 25, wherein the promoter is			
1 2	selected from	the gr	oup consisting of a tetracycline responsive promoter, a probasin			
3 🗇	promoter, an	d a CM	IV promoter.			
<b>j.</b> .7			•			
1	/	36.	A method of transducing a cell with a nucleic acid encoding a			
2 1	viral inhibitor	r comp	rising contacting the cell with the cell transduction vector of claim			
3 <u>1</u> 4	1.	•				
	. )	_37.	The method of claim 36, wherein the cell is transduced in vitro.			
	$W_{1}/\sqrt{2}$	7"	The method of claim 50, wherein the con is transdaced to via or			
. //	A /	70	A week of a finite in the arounth of HIV in a call comprising			
1	V'	38.	A method of inhibiting the growth of HIV in a cell comprising			
2	transducing t	ne cell	with the cell transduction vector of claim 1.			
1		39.	The method of claim 38, wherein the cell is isolated from a			
2	mammal, and wherein the method further comprises introducing the cell into a					
3	mammal.					
$^{(1)}$		40.	The method of claim 39, wherein the cell is selected from the			
2/1/2-	group of cell	s consi	sting of transferrin receptor+ cells, CD4+ cells and CD34+			
	hematopoieti	;	·			
" U" /	, -					
1	0151>	41.	A cell comprising the cell transduction vector of claim 1.			

The cell of claim 41, wherein the cell is selected from the group of cells comprising CD4+ cells, CD34+ hematopoietic stem cells, and transferrin receptor+ cells.